```
L11 ANSWER 1 OF 8 USPATFULL on STN
       1998:1990 USPATFULL
AN
       Integrated process for producing disopropyl ether, an isopropyl
TI
       tertiary alkyl ether and isopropyl alcohol
       Frey, Stanley J., Palatine, IL, United States
Schmidt, Robert J., Barrington, IL, United States
Marker, Terry L., Warrenville, IL, United States
IN
       Marinangeli, Richard E., Arlington Heights, IL, United States
       UOP, Des Plaines, IL, United States (U.S. corporation)
PΑ
PΙ
       US 5705712
                                19980106
ΑI
       US 1995-539394
                                19951005 (8)
DT
       Utility
       Granted
FS
       Primary Examiner: Cintins, Marianne M.; Assistant Examiner: Jones,
EXNAM
       Dwayne C.
       McBride, Thomas K., Snyder, Eugene I., Maas, Maryann
LREP
CLMN
       Number of Claims: 14
       Exemplary Claim: 1
ECL
DRWN
       2 Drawing Figure(s); 2 Drawing Page(s)
LN.CNT 630
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       A highly integrated process for concurrently producing diisopropyl ether
       and an isopropyl tertiary alkyl ether has been developed. Optionally,
       high purity isopropyl alcohol may also be collected as a product. In a
       first reactor, propylene and water are reacted to form isopropyl
       alcohol, a portion of which is further reacted to form diisopropyl
       ether. After removing unreacted propylene, the effluent of the first
       reactor is separated into an ether rich stream, a water rich
       stream and an alcohol rich stream. The alcohol rich
       stream is dried to provide dry isopropyl alcohol. A portion of
       the dry isopropyl alcohol may be removed and collected as a product. A
       portion of the dry isopropyl alcohol and isobutylene, isoamylene or a
       mixture thereof are reacted to form an isopropyl tertiary alkyl ether in
       a second reactor. Unreacted iso-olefins and inert compounds are then
       removed from the second reactor effluent. A mixture of the effluent from
       the second reactor and the ether rich and the water rich streams
       separated from the first reactor are water washed to produce a mixed
       ethers product stream and an aqueous isopropyl alcohol recycle
       stream. The isopropyl tertiary alkyl ether is collected along
       with the diisopropyl ether in the mixed ethers product stream
       from the water wash. A modified flowscheme of the process is also
       discussed.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 2 OF 8 USPATFULL on STN
L11
       97:107294 USPATFULL
AN
       Integrated process for producing diisopropyl ether and an isopropyl
ΤI
       tertiary alkyl ether
       Frey, Stanley J., Palatine, IL, United States
IN
       Schmidt, Robert J., Barrington, IL, United States
       Marker, Terry L., Warrenville, IL, United States
       Marinangeli, Richard E., Arlington Heights, IL, United States
       UOP, Des Plaines, IL, United States (U.S. corporation)
PΑ
                                19971118
PΙ
       US 5689014
ΑI
       US 1995-539577
                                19951005 (8)
       Utility
DT
FS
       Granted
       Primary Examiner: Cintins, Marianne M.; Assistant Examiner: Jones,
EXNAM
       Dwayne C.
       McBride, Thomas K., Snyder, Eugene I., Maas, Maryann
LREP
CLMN
       Number of Claims: 14
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ECL

Exemplary Claim: 1

1 Drawing Figure(s); 1 Drawing Page(s) DRWN LN.CNT 534 CAS INDEXING IS AVAILABLE FOR THIS PATENT. A highly integrated process for concurrently producing diisopropyl ether and an isopropyl tertiary alkyl ether has been developed. In a first reactor, propylene and water are reacted to form isopropyl alcohol, a portion of which is further reacted to form diisopropyl ether. After removing unreacted propylene, the effluent of the first reactor is separated into an ether rich stream, a water rich stream and an alcohol rich stream. The alcohol rich stream and isobutylene, isoamylene or a mixture thereof are reacted to form an isopropyl tertiary alkyl ether in a second reactor. The water present in the alcohol rich stream also reacts with the iso-olefin to form tertiary alcohol. The effluent from the second reactor is water washed to produce an oxygenate product stream and an aqueous alcohol recycle stream. Some tertiary alcohol is recycled to the first reactor where it is reacted with propylene to form additional isopropyl tertiary alkyl ether. The isopropyl tertiary alkyl ether and some tertiary alcohol is collected along with the diisopropyl ether in the mixed oxygenate product stream from the water wash. CAS INDEXING IS AVAILABLE FOR THIS PATENT. ANSWER 3 OF 8 USPATFULL on STN L11 97:78233 USPATFULL AN Process for producing slippery, tenaciously adhering hydrogel coatings containing a polyurethane-urea polymer hydrogel commingled with a poly (n-vinylpyrrolidone) polymer hydrogel Hostettler, Fritz, Lambertville, NJ, United States IN Rhum, David, New York, NY, United States Forman, Michael R., Ramsey, CA, United States Helmus, Michael N., St. Louis Park, MN, United States Ding, Ni, Plymouth, MN, United States Schneider (USA) Inc., Plymouth, MN, United States (U.S. corporation) PA 19970902 PΤ US 5662960 ΑI US 1995-384711 19950201 (8) Utility DT Granted FS Primary Examiner: Dudash, Diana EXNAM Richardson, Peter C., Akers, Lawrence C., Jaeger, Howard R. LREP CLMN Number of Claims: 80 ECL Exemplary Claim: 1 DRWN No Drawings LN.CNT 3423 CAS INDEXING IS AVAILABLE FOR THIS PATENT. A process for preparing coating compositions of a commingled hydrogel of AB a polyurethane-polyurea polymer hydrogel and a poly(N-vinylpyrrolidone) polymer hydrogel; a process for making materials composed of a polymeric plastic or rubber substrate or a metallic substrate, with a coating of the commingled hydrogel thereon; and a process for making medical devices with a coating of the commingled hydrogel thereon, are disclosed. The coating compositions tenaciously adhere to the substrate materials and medical devices to which they are applied due to bonding of a tie coat to a reactive substrate surface and due to the commingling

of the two hydrogel components. The coating compositions and coated materials and medical devices are non-toxic and biocompatible, making them ideally suited for use in applications such as for catheters, catheter balloons and stents. In such applications, the coating compositions, coated materials, and coated medical devices made

therefrom demonstrate low coefficients of friction in contact with body fluids, especially blood, as well as a high degree of wear permanence over prolonged use. The commingled hydrogel coatings are capable of being dried to facilitate storage of the devices to which they have been applied, and can be instantly reactivated for later use by exposure to water.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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L11 ANSWER 4 OF 8 USPATFULL on STN
AN
       94:55683 USPATFULL
       Integrated process for producing diisopropyl ether from isopropyl
ΤI
       alcohol
       Marker, Terry L., Warrenville, IL, United States
IN
       Kempf, Laura E., Deerfield, IL, United States
PA
       UOP, Des Plaines, IL, United States (U.S. corporation)
PΙ
       US 5324866
                               19940628
                               19930323 (8)
       US 1993-36008
AΙ
       Utility
DT
       Granted
FS
EXNAM Primary Examiner: Mars, Howard T.
       McBride, Thomas K., Spears, Jr., John F., Taylor, Reginald K.
LREP
       Number of Claims: 16
CLMN
       Exemplary Claim: 1
ECL
       1 Drawing Figure(s); 1 Drawing Page(s)
DRWN
LN.CNT 587
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The present invention is an integrated isopropyl alcohol (IPA) and
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The present invention is an integrated isopropyl alcohol (IPA) and diisopropyl ether (DIPE) process. In this process, IPA, substantially free of DIPE, is formed in a hydration reactor by reacting an olefinic feedstock with water in a hydration reactor. The effluent from the hydration reactor is then contacted in a first separation unit with DIPE which was made in an etherification reactor. The resulting mixture is then passed to a second separation unit to separate the IPA from the DIPE product. The IPA is then fed to the etherification reactor to produce DIPE.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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L11 ANSWER 5 OF 8 USPATFULL on STN
       94:55682 USPATFULL
AN
       Di-isopropyl ether production
TI
       Beech, Jr., James H., Wilmington, DE, United States
IN
       Miller, Douglas, Yardley, PA, United States
       Soto, Jorge L., Cranbury, NJ, United States
       Stoos, James A., Blackwood, NJ, United States Wu, Albert H., Medford, NJ, United States
       Mobil Oil Corporation, Fairfax, VA, United States (U.S. corporation)
PA
PΙ
       US 5324865
                                19940628
ΑI
       US 1993-20964
                                19930222 (8)
DT
       Utility
       Granted
FS
       Primary Examiner: Mars, Howard T.
EXNAM
       McKillop, Alexander J., Keen, Malcolm D., Wise, L. Gene
LREP
       Number of Claims: 14
CLMN
ECL
       Exemplary Claim: 1
       2 Drawing Figure(s); 2 Drawing Page(s)
DRWN
LN.CNT 753
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       A process for production of diisopropyl ether by conversion of
       hydrocarbon feedstock containing propene, propane and C.sub.2 - light
       gas components, including the steps of: optionally, prefractionating
       fresh feedstock containing propene, propane and C.sub.2 - light gas
       components to provide a reactor feedstream rich in propene; contacting
       the feedstock and water in a catalytic reactor with acidic
       catalyst under olefin hydration and etherification conditions;
       and recovering from the catalytic reactor a liquid reactor
       effluent stream containing diisopropyl ether, isopropanol,
       water, unreacted propene, propane and C.sub.2 - light gas components.
       Improved operation is achieved by separating the liquid effluent
       stream in a vertical stripper column; recovering an
       overhead vapor stream containing propene, propane and C.sub.2
       - light gas components from the stripper column; cooling the
       overhead vapor stream to provide a reflux stream
       rich in condensed propene and propane; removing the C.sub.2 - light gas
       components from condensed; recycling the reflux stream to an
       upper contact portion of the stripper column; and recovering a
       predominantly C3 recycle stream from the upper contact portion
       of the stripper column. Optionally, the C3 recycle
       stream may be passed to the to the prefractioning step for
       propene enrichment with fresh feedstock.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 6 OF 8 USPATFULL on STN
L11
       93:102927 USPATFULL
AN
       Process of preparing an isopropanol and diisopropyl ether oxgenate motor
ΤI
       fuel additive
       Irvine, Robert L., Overland Park, KS, United States
IN
       The Pritchard Corporation, Overland Park, KS, United States (U.S.
PA
       corporation)
PΤ
       US 5268515
                                19931207
                                19930301 (8)
       US 1993-24525
AΤ
       Continuation-in-part of Ser. No. US 1992-877642, filed on 1 May 1992,
RLI
       now patented, Pat. No. US 5191129
DТ
       Utility
FS
       Granted
       Primary Examiner: Lone, Werren B.
EXNAM
       Hovey, Williams, Timmons & Collins
LREP
CLMN
       Number of Claims: 14
ECL
       Exemplary Claim: 1
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DRWN

7 Drawing Figure(s); 7 Drawing Page(s)

LN.CNT 1993

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Propene is reacted with water in a multi-stage, fluidized bed catalytic reactor to produce an oxygenate motor fuel additive containing a major proportion of isopropanol, a minor proportion of diisopropyl ether, and some water. The molar ratio of water to propene introduced into each catalytic stage of the multi-stage reactor is maintained within a range of from about 2:1 to about 6:1. The temperature of the reactants in each of the catalytic stages is maintained within a range of from about 250° F. to about 320° F. and the pressure at a level of from about 1200 psia to about 3600 psia. The temperature in each catalytic stage increases from the initial catalytic stage to the final catalytic stage with the temperature increase being limited to a value within a range of from about 8° F. to about 1° F. The pressure of the final stage of the catalytic section is controlled so that the reaction product containing the organic constituents including oxygenates provides a concentrated, less dense liquid stream which may be easily separated from the aqueous liquid phase in the final stage. The liquid phase is recycled to the catalytic zone while underacted propene is separated from the oxygenates. Some water is purposefully retained in the oxygenate to take advantage of the solubilization of the IPA component. Seasonal gasoline component specifications may be met by simply controlling the amount of water allowed to remain in the oxygenate product.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L11 ANSWER 7 OF 8 USPATFULL on STN

AN 93:16845 USPATFULL

TI Method of preparing an isopropanol and diisopropyl ether oxygenate motor fuel additive

IN Irvine, Robert L., Overland Park, KS, United States

PA The Pritchard Corporation, Overland Park, KS, United States (U.S.

corporation)

PI US 5191129 19930302

AI US 1992-877642 19920501 (7)

DT Utility

FS Granted

EXNAM Primary Examiner: Lone, Werren B.

LREP Hovey, Williams, Timmons & Collins

CLMN Number of Claims: 22

ECL Exemplary Claim: 1

DRWN 3 Drawing Figure(s); 3 Drawing Page(s)

LN.CNT 1245

AB

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Propene is reacted with water in a multi-stage, fluidized bed catalytic reactor to produce an oxygenate motor fuel additive containing a major proportion of isopropanol (IPA), some diisopropyl ether (IPE) and some water. The molar ratio of water to propene introduced into each catalytic stage of the multi-stage reactor is maintained within a range of from about 2:1 to about 6:1. The temperature of the reactants in each of the catalytic stages is maintained within a range of from about 250° F. to about 300° F. and the pressure at a level of from about 1200 psia to about 3000 psia. The temperature in each catalytic stage increases from the initial catalytic stage to the final catalytic stage with the temperature increase being limited to a value within a range of from about 8° F. to about 2° F. The pressure of the final stage of the catalytic section is controlled so that the reaction product containing the organic constituents including oxygenates provides a concentrated, less dense liquid stream which may be easily separated from the aqueous liquid phase in the final stage. The liquid phase is recycled to the

catalytic zone while unreacted propene is separated from the oxygenates. Some water is purposefully retained in the oxygenate to take advantage of the solubilization of the IPA component. Seasonal gasoline component specifications may be met by simply controlling the amount of water allowed to remain in the oxygenate product. Relatively small distillation towers are suitable in the present process because of the 92% conversion factor of the propene entering. Thus, only about 1/3rd of the quantity of unconverted hydrocarbons must be separated in the present process as compared with existing commercial methods for the same yield of motor fuel oxygenates.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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L11 ANSWER 8 OF 8 USPATFULL on STN
       88:50083 USPATFULL
AN
       Isopropyl alcohol purification process
TI
       Litzen, David B., Houston, TX, United States
IN
       Bolger, Stephen R., La Porte, TX, United States
       Shell Oil Company, Houston, TX, United States (U.S. corporation)
PΑ
                               19880809
PΙ
       US 4762616
      US 1986-943356
                               19861219 (6)
ΑI
DT
      Utility
FS
      Granted
EXNAM Primary Examiner: Spear, Frank
      Lemuth, Richard F.
LREP
      Number of Claims: 9
CLMN
      Exemplary Claim: 1
ECL
       1 Drawing Figure(s); 1 Drawing Page(s)
DRWN
LN.CNT 476
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       A process for the purification of the crude isopropyl alcohol product of
AB
       propylene hydration reactions. Crude isopropyl alcohol products, which
       comprise isopropyl alcohol, diisopropyl ether and polymeric impurities,
       are subjected to a specified sequence of multiple dilution and phase
       separation steps which serve to extract a substantial portion of the
       product's diisopropyl ether and polymeric impurities. The invention is
       particularly useful in removing odiferous sulfur-containing impurities
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from the products of indirect propylene hydration processes which involve the reaction of propylene with sulfuric acid to produce isopropyl sulfate followed by hydrolysis of the sulfate to

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

isopropyl alcohol.

ANSWER 6 OF 7 USPATFULL on STN L1397:49823 USPATFULL ΑN Isopropyl alcohol and diispropyl ether production from crude by-product ΤI acetone in one step Taylor, Jr., Robert J., Port Arthur, TX, United States INDai, Pei-Shing E., Port Arthur, TX, United States Knifton, John F., Austin, TX, United States Martin, Bobby R., Beaumont, TX, United States Texaco Chemical Inc., White Plains, NY, United States (U.S. corporation) PAUS 5637778 19970610 PΙ US 1994-287451 19940808 (8) AIUtility DТ FS Granted Primary Examiner: Dees, Jose'G.; Assistant Examiner: Williams, Rosalynd **EXNAM** Bailey, James L., Priem, Kenneth R., Hunter, Cynthia L. LREP Number of Claims: 17 CLMN ECL Exemplary Claim: 1 1 Drawing Figure(s); 1 Drawing Page(s) DRWN LN.CNT 786 CAS INDEXING IS AVAILABLE FOR THIS PATENT. Disclosed is a one-step method for synthesis of methyl tertiary butyl ether, diisopropyl ether and isopropyl ether from crude streams containing acetone, methanol and t-butyl alcohol which comprises reacting an acetone-rich feed over a bifunctional catalyst

- Disclosed is a one-step method for synthesis of methyl tertiary butyl ether, diisopropyl ether and isopropyl ether from crude streams containing acetone, methanol and t-butyl alcohol which comprises reacting an acetone-rich feed over a bifunctional catalyst comprising 5-45% by weight of a catalyst consisting essentially of a hydrogenation catalyst selected from the group consisting of one or more metals selected from the group consisting of IB, VIB or VIII of the Periodic Table and a heteropoly acid on a 55%-95% of the total weight of the catalyst of a support comprising a compound selected from the group consisting of:
 - a. a metal phosphate;
 - b. 5 to 95% by weight metal phosphate supported on 95 to 5 wt % Group III or IV oxide; and
 - c. a large pore silicoaluminophosphate.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

(FILE 'HOME' ENTERED AT 18:57:41 ON 10 MAY 2004)

L1 L2	FILE 'REGISTRY' ENTERED AT 18:58:01 ON 10 MAY 2004 1 S DIISOPROPYL ETHER/CN 1 S ISOPROPANOL/CN
	FILE 'CAPLUS, USPATFULL, CA, CAOLD' ENTERED AT 18:59:10 ON 10 MAY 2004
L3	7387 S L1
L4	2103 S L1 AND L2
L5	274 S L4 AND DISTILL?
L6	145 S L5 AND CATALY?
L7	37 S L6 AND HYDROLY?
L8	15 S L7 AND COLUMN
L9	8 S L8 AND RESIN
L10	8 DUP REM L9 (0 DUPLICATES REMOVED)
L11	8 S L10 AND STREAM
L12	7 S L8 NOT L11
L13	7 DUP REM L12 (0 DUPLICATES REMOVED)